

wherein:

R<sub>1</sub> and R<sub>2</sub> are independently selected from hydrogen and alkyl;

R<sub>3</sub> is alkyl;

B<sub>1</sub>  
ant  
R<sub>4</sub>, R<sub>6</sub> and R<sub>7</sub> are independently selected from hydrogen, halogen, hydroxy, alkyl, aryl, amino, alkylamino, dialkylamino, alkoxy, aryloxy, alkylthio, alkylsulfoxyl, alkylsulfonyl, nitro, carbonitrile, carbo-alkoxy, carbo-aryloxy and carboxyl;

R<sub>5</sub> is selected from hydrogen, halogen, hydroxy, alkyl, aryl, amino, alkylamino, dialkylamino, alkoxy, aryloxy, alkylthio, alkylsulfoxyl, alkylsulfonyl, nitro, carbonitrile, carbo-alkoxy, carbo-aryloxy and carboxyl; and

A is a 5-membered partially unsaturated or aromatic heterocyclic ring or a 5-membered partially unsaturated carbocyclic ring,

or a pharmaceutically acceptable salt, addition compound or prodrug thereof.

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B<sub>2</sub>  
13. (Amended Twice) A compound according to claim 1, wherein A is a 5-membered partially unsaturated carbocyclic ring or a 5-membered partially unsaturated or aromatic heterocyclic ring.

14. (Amended Twice) A compound according to claim 1, wherein A is selected from cyclopentenyl and thienyl.

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